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Plasmodium falcipa	6 AAQ80908	16	2223	1.3	48.8	45	
cDNA encoding Plas	3 AAQ29187	13	2223	1.3	48.8	44	
Staphylococcus aur	8 AAV74833	16	1987	1.3	48.8	4 .	O
P. falciparum live	8 AAT78867	18	6152	1.3	49	42	
P. falciparum live		18	5361	1.3	49	41	
Sequence encoding	1 AAQ03875	11	3095	1.4	50	40	
Numan dentin sialo	1 AAA88664	21	8201	1.4	50.2	39	O
Plasmodium falcipa		21	1998	1.4	50.6	38	
Plasmodium falcipa	1 AAA70095	2	7326	1.4	50.8	37	
Staphylococcus aur	8 AAT91077	18	2367	1.4	51	36	
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Haemophilus adhesi	7 AAT41476	17	7291	1.4	51.4	34	
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Borrelia burgdorfe	0 AAX20252	20	35515	1.4	51.8	32	O
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S. aureus RUSA266		2	4248	1.5	56.8	24	
Borrelia burgdorfe	0 AAX20280	20	3653	1.6		23	
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DNA encoding Leuco	6 AAQ87587	16	1686	1.7	62.4	21	
Plasmodlum falcipa	1 AAA70099	2	3579	1.7	. 62.6	20	
Chicken leucocytoz		17	3399	2.1	77.4	19	
Oligonucleotide Dl		N N	938	4.0	147.2	18	O
		22	936	4.0	147.2	17	O
	2 AAF58259	2	936	4.0	147.2	16	O
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Oligonucleotide Dl	2 AAF58254	Ņ	9E6	4.0	147.2	14	O
	2 AAF58252	22	936	4.0	147.2	13	c
Oligonucleotide D1		N	958	4.1	150.6		

ALIGNMENTS

RESULT AAT75487

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                        01-FEB-1996;
                                 07-AUG-1997.
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S-layer; sbs-A; vaccine; adjuvant; carrier; hybridisation assay; molecular spinning nozzle: molecular laser; ds.

DNA for Bacillus stearothermophilus S-layer protein sbs-A

27-FEB-1998 (first entry)

AAT75487;

AAT75487 standard; DNA; 3687

BP

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Query Match
Best Local
Matches 368
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Preparation of S-layer proteins by expressing negative bacterlum - or new sbs-B gene in any recombinant proteins containing heterologous i epitope(s), useful as vaccines and adjuvants
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AAX22748 standard; DNA; 3687 쁌

24-AUG-1999 entry)

stearothermophilus sbsA DNA.

sbsA protein; integration; eukaryotic ho; protein; cytop1 S-layer toplasmic cell; vac protein; vaccine; membrane; adjuvant; Gram-negative; prokaryotic ane; secretion; periplasmic immunogenic epitope; host cell: luciferase;

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                                                                                                                                     This invention describes a method for the production of a S-layer protein CC (I) which comprises (a) preparing a Gram-negative prokaryotic host cell ctransformed with nucleic acid (II) encoding (I), linked to a signal CC sequence (SS) that encodes a protein which causes at least one of (1) CC integration of (I) into the external or cytoplasmic membranes and/or (1) into the periplasmic space or extracellular medium, CC (ii) secretion of (I) into the periplasmic space on extracellular medium, CC (ii) the membranes, periplasmic space and/or extracellular medium, CC (ii) culturing the cell to express (I) and (C) optionally recovering (I) CC from the membranes, periplasmic space and/or extracellular medium, CC (ii) and (ii) a secretion into the extracellular medium, (I), and (iii) and (iii) as interested structures, may include a wide variety of polypeptide lnserts and CC organelle and/or secretion into the extracellular medium, (I), and (iii) as useful as (I) vaccines or adjuvants (with immunogenic epitopes or Immunostinulants inserts such as cytokines) (I) as reactors (inserts CC are enzymes, e.g. polyhydroxybutyrate (PHB) synthase for use as a CC (insert and inserts such as cytokines) (I) as reactors (inserts CC are enzymes, e.g. polyhydroxybutyrate (PHB) synthase for use as conjuvant laser (when combined with substrate and oxygen)) and (iii) as conjuvant carrier molecule (streptavidin is inserted) for use in CC (inserts) in this system, heterologous (I) do not form inclusion bodies but rather monomolecular layers, and in eukaryotic cells they undergo CC glycosylation. This sequence encodes the Bacillus stearothermophilus stearothermophilus sond to invention.
                                                                       Query Match
Best Local Similarity
Matches 3687; Conser
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            producing S-layer proteins in Gram-negative bacteria or eukaryotes integrated into membranes or organelies or secreted into periplasme or growth medium, and nucleic acid encoding S-layer proteins with peptide insertions, used in vaccines or for enzymatic reactions
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1800	gaaaatacaatctatcaaatcaaaattaaaaaaggcttgaagtccgataaaggtattgaa 	1741 1741	p 99
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polyhydroxyalkanoate synthesis;
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describes a novel host cell (A) comprising at least ombinant polypeptides (I), at least one being in carr e products of the invention have antibacterial activity
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Best Loc
Matches
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body fluid;
yer; Gene 3;
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This invention describes a method for the production of a S-layer protein CC (I) which comprises (a) preparing a Gram-negative prokaryotic host cell transformed with nucleic acid (II) encoding (I), linked to a signal CC sequence (SS) that encodes a protein which causes at least one of (i) (ii) secretion of (I) into the external or cytoplasmic membranes and/or CC (ii) secretion of (I) into the periplasmic space or extracellular medium, (CC integration of (I) into the periplasmic space or extracellular medium, (CC integratively, a eukaryotic cell is used as host and then the SS, which (CC is optional) promotes integration into the cytoplasmic membrane or an organelle and/or secretion into the extracellular medium. (I), and (CC derived structures, may include a wide variety of polypeptide inserts and care useful as (i) vaccines or adjuvants (with immunogenic epitopes or communostimulants inserts such as cytokines) (ii) as reactors (inserts are enzymes, e.g. polyhydroxybutyrate (PHB) synthase for use as a conjuvarial laser (when combined with substrate and oxygen)) and (iii) as universal carrier molecule (streptavidin is inserted) for use in the cytopes). In this system, heterologous (I) do not form inclusion bodies out rather monomolecular layers, and in eukaryotic cells they undergo constructed from the Bactllus stearothermophilus sbsA gene and the Bacteriophage fd Gene 3 signal sequence which is used in the method of the invention.

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stearothermophilus sbsA/malE fusion protein UNA sequence

entry)

24-AUG-1999 AAX78245;

sbsA protein; S-layer protein; Gram-negative; prokaryotic host or integration: cytoplasmic membrane; secretion; periplasmic space; eukaryotic host cell; vaccine; adjuvant; immunogenic epitope; luminostimulant; cytokine; polyhydroxybutyrate; PHB synthase; box molecular laser; universal carrier molecule; monomolecular layer. protein; malE; synthase; body lecular layer; toxin,

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S

Producing S-layer proteins in Gram-negative bacteria or eukaryotes integrated into membranes or organelles or secreted into periplasm or growth medium, and nucleic acid encoding S-layer proteins with peptide insertions, used in vaccines or for enzymatic reactions eukaryotes

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This invention describes a method for the production of a S-layer protei (I) which comprises (a) preparing a Gram-negative prokaryotic host cell transformed with nucleic acid (II) encoding (I), linked to a signal encodes a protein аt (i)

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integration of (I) into the external or cytoplasmic membranes and/or (ii) secretion of (I) into the periplasmic space or extracellular medium, (C) cytoplasmic space and/or extracellular medium, (I) cytoplasmic space and/or extracellular medium, (I) cytoplasmic space and/or extracellular medium. (I) cytoplasmic space and/or extracellular medium. (I) and (c) optionally recovering (I) cytoplasmic space and/or extracellular medium. (I) cytoplasmic membrane or an (I) and (I) extracellular medium. (I), and (I) and (I) extracellular medium. (I), and (I) are useful as (I) vaccines or adjuvants (with immunogenic epitopes or (I) are useful as (I) vaccines or adjuvants (with immunogenic epitopes or (I) are enzymes, e.g. polyhydroxybutyrate (PHB) synthase for use as a (I) vaccines as a cytokines) (II) as reactors (inserts as cytokines) (II) as cytoplasmic epitopes) as (II) as cytoplasmic epitopes) and (III) as (Inserts are enzymes, e.g. polyhydroxybutyrate (PHB) synthase for use in (I) expressed carrier molecule (streptavidin is inserted) for use in (I) expressed carrier molecule (streptavidin is inserted) for use in (I) epitopes). In this system, heterologous (I) do not form inclusion bodies (I) cytokines, toxins etc. from body fluids (inserts are specific binding (I) cytopidisation and immuno assays, or for selective elimination of (I) cytokines, toxins etc. from body fluids (inserts are specific binding (I) cytopidisation. This sequence encodes a novel fusion protein constructed (I) cytopidisation. This sequence encodes a novel fusion protein constructed (I) esquence which is used to illustrate the method of the invention.
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different redox potentials. The invention is used for electronic
detection of nucleic acids, especially of substitutions (mismatches)
and single-nucleotide polymorphisms, e.g. for genotyping,
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1.18; Pred. No. 1.2e
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                                                                                                                                                                                                                                  acids each containing an electron-transfer group (ETM) having different redox potentials. The invention is used for electronic detection of nucleic acids, especially of substitutions (mismatches) and single-nucleotide polymorphisms, e.g. for genotyping,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Electron-transfer group; gene expression; ss.
124
                                                                                                                                                                                                   Sequence 936 BP; 6 A; 138 C; 8 G; 8
                                                                                                                                                                                                                                                                         The present invention relates to a composition comprising two nucleic acids each containing an electron-transfer group (ETM) having
                                                                                                                                                                                                                                                                                                                                    Nucleic acids containing electron-transfer group, useful as labels in hybridization assays, e.g. for genotyping, allowing repeat analyses of a single surface.
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17-MAR-2000; 2000US-0190259
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                                                                                                                                        / Match 4.1%; Score 150.6; DB 22;
Local Similarity 1.1%; Pred. No. 1.2e-23;
hes 9; Conservative 500; Mismatches 272;
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Nucleic acids containing electron-transfer group, useful as labels in hybridization assays, e.g. for genotyping, allowing repeat analyses on a single surface
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17-MAR-2000;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Nucleic acids containing electron-transfer group, useful as labeis in hybridization assays, e.g. for genotyping, allowing repeat analyses on
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The present invention relates to a composition comprising two nuclei acids each containing an electron-transfer group (ETM) having different redox potentials. The invention is used for electronic detection of nucleic acids, especially of substitutions (mismatches) and single-nucleotide polymorphisms, e.g. for genotyping,

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